

Supp. Data Table S2 Investigated genes and associated diseases

Disease	Gene	CHR	Protein name	RefSeq-Nr.	other associated diseases	
<b>Cardiomyopathies</b>						
<b>ARVC</b>	<i>CTNNA3</i>	10	Catenin (cadherin-associated protein), alpha 3	NM_001127384.2		
	<i>DSC2</i>	18	Desmocollin 2	NM_024422.4		
	<i>DSG2</i>	18	Desmoglein 2	NM_001943.3	DCM	
	<i>DSP</i>	6	Desmoplakin	NM_004415.2	DCM	
	<i>JUP</i>	17	Junction plakoglobin	NM_021991.2		
	<i>PKP2</i>	12	Plakophilin2	NM_004572.3	BrS	
	<i>TGFβ3</i>	14	Transforming growth factor B3	NM_003239.2	LDS, TAAD	
	<i>TMEM43</i>	3	Transmembrane protein 43	NM_024334.2	EDMD	
<b>DCM</b>	<i>ABCC9</i>	12	ATP-binding cassette, subfamily C (CFTR/MRP), member 9	NM_005691.3	Afib, BrS	
	<i>ACTN2</i>	1	Actinin, alpha 2	NM_001103.3	HCM	
	<i>BAG3</i>	10	Bcl2-associated athanogene 3	NM_004281.3	muscular dystrophy	
	<i>CTF1</i>	16	Cardiotrophin 1	NM_001330.3	hypertensive heart disease	
	<i>CRYAB</i>	11	Crystallin, a B	NM_001289807.1	fatal infantile hypertonic myofibrillar myopathy	
	<i>DES</i>	2	Desmin	NM_001927.3	limb-girdle muscular dystrophy	
	<i>DMD</i>	X	Dystrophin, muscular dystrophy	NM_004006.2	duchenne muscular dystrophy, becker muscular dystrophy	
	<i>DNAJC19</i>	3	DNAJ (Hsp40) homolog, subfamily C, member 19	NM_145261.3	3-methylglutaconic aciduria	
	<i>DOLK</i>	9	Dolichol kinase	NM_014908.3	congenital disorder of glycosylation	
	<i>EYA4</i>	6	Eyes absent homolog 4 (Drosophila)	NM_172105.3	nonsyndromic hearing loss and deafness	
	<i>FHL2</i>	2	Four and a half LIM domains 2	NM_001450.3	hemophagocytic lymphohistiocytosis	
	<i>FKTN</i>	9	Fukuyama-type congenital muscular dystrophy (fukutin)1	NM_006731.2	muscular dystrophy-dystroglycanopathy	
	<i>GAA</i>	17	Glucosidase, alpha acid	NM_000152.3	congenital disorder of glycosylation	
	<i>GATAD1</i>	7	GATA zinc finger domain containing 1	NM_021167.4		
	<i>ILK</i>	11	Integrin-linked kinase	NM_001014794.2		
	<i>LAMA4</i>	6	Laminin, alpha 4	NM_001105206.2		
	<i>LDB3</i>	10	LIM binding domain 3 (ZASP)	NM_001080114.1	HCM, LVNC	
	<i>LMNA</i>	1	Lamin A/C	NM_170707.2	ARVC, EDMD	
	<i>MYPN</i>	10	Myopalladin	NM_001256267.1	HCM	
	<i>NEBL</i>	10	Nebulette	NM_006393.2		
	<i>NEXN</i>	1	Nexilin (F actin-binding protein)	NM_144573.3	HCM, ASD, LVNC	
	<i>PLN</i>	6	Phospholamban	NM_002667.3	HCM	
	<i>PSEN1</i>	14	Presenilin-1	NM_000021.3		
	<i>PSEN2</i>	1	Presenilin-2	NM_000447.2		
	<i>RBM20</i>	10	RNA-binding motif protein 20	NM_001134363.2		
	<i>RAF1</i>	3	Raf-1 proto-oncogene, serine/threonine kinase	NM_002880.3	Noonan syndrome	
	<i>SDHA</i>	5	Succinate dehydrogenase complex, subunit A, flavoprotein (Fp)	NM_001294332.1	mitochondrial complex deficiency	
	<i>SGCD</i>	5	Sarcoglycan delta (ystrophin Ass glycoprotein	NM_000337.5	lim-girdle muscular dystrophy	
	<i>TAZ</i>	X	Tafazzin	NM_000116.3	LVNC, BTHS	
	<i>TCAP</i>	17	Titin cap (telethonin)	NM_003673.3	HCM, limb-girdle muscular dystrophy	
	<i>TMPO</i>	12	Thymopoietin	NM_003276.2	muscular dystrophy	
	<i>TNNC1</i>	3	Troponin C type 1	NM_003280.2	HCM	
	<i>TNN3</i>	19	Troponin I Type 3 (cardiac)	NM_000363.4	HCM, Afib	
	<i>TPM1</i>	15	Tropomyosin 1 alpha	NM_000366.5		
	<i>TRPM7</i>	15	Transient receptor potential cation channel, subfamily M, member 7	NM_001301212.1	amyotrophic lateral sclerosis-parkinsonism/dementia complex	
	<i>TSMF</i>	12	Ts translation elongation factor, mitochondrial	NM_001172695.1	combined oxidative phosphorylation deficiency, fatal mitochondrial disease	
	<i>TTR</i>	18	Transthyretin	NM_000371.3	hyperthyroxinemia, dysransthyretinemic	
	<i>TXNRD2</i>	22	Thioredoxin reductase 2	NM_006440.4		
	<i>XK</i>	X	x-linked Kx blood group (McLeod syndrome)	NM_021083.2		
	<i>ZASP</i>	7	ZO-2 associated speckle protein	NM_001289933.1	HCM, myofibrillar myopathy	
	<b>HCM</b>	<i>ACTC1</i>	15	Actin, alpha, cardiac muscle 1	NM_005159.4	ASD, DCM, LVNC
		<i>ANKRD1</i>	10	Ankyrin repeat domain 1 (cardiac muscle)	NM_014391.2	DCM
<i>CALR3</i>		19	Calreticulin 3	NM_145046.4		
<i>CAV3</i>		3	Caveolin 3	NM_001234.4	LQTS, limb-girdle muscular dystrophy	
<i>CSRP3</i>		11	Cysteine- and glycine-rich protein 3 (cardiac LIM protein)	NM_003476.4	DCM	
<i>JPH2</i>		20	Junctophilin 2	NM_020433.4	Afib	
<i>LAMP2</i>		X	Lysosome associate membrane glycoprotein 2	NM_001122606.1	GSD	
<i>MAP2K1</i>		15	Mitogen-activated protein kinase kinase 1	NM_002755.3		
<i>MAP2K2</i>		19	mitogen-activated protein kinase kinase 2	NM_030662.3	neurofibromatosis-noonan syndrome	
<i>MRPL3</i>		3	mitochondrial ribosomal protein L3	NM_007208.3	combined oxidative phosphorylation deficiency	
<i>MYBPC3</i>		11	Myosin binding protein C, cardiac	NM_000256.3	DCM, LVC	
<i>MYH6</i>		14	Myosin, heavy chain, cardiac a	NM_002471.3	ASD, DCM	
<i>MYH7</i>		14	Myosin Heavy chain 7, cardiac B	NM_000257.3	DCM, myosin storage myopathy	
<i>MYL2</i>		12	Myosin light chain 2, regulatory, cardiac	NM_000432.3	DCM	
<i>MYH11</i>		16	Myosin heavy chain 11, smooth muscle	NM_001040114.1	TAAD,	
<i>MYL3</i>		3	Myosin light chain 3, ventricular, skeletal, slow	NM_000258.2	DCM	
<i>MYLK</i>		3	Myosin light chain kinase	NM_053025.3	TAAD,	
<i>MYLK2</i>		20	Myosin light chain kinase 2	NM_033118.3		
<i>MYOZ2</i>		4	Myozenin 2	NM_016599.4	DCM	
<i>MYOM1</i>		18	Myomesin 1	NM_003803.3		
<i>PRKAG2</i>		7	Protein kinase, AMP-activated, 2 noncatalytic subunit	NM_016203.3	glycogen storage disease of heart	
<i>PDLIM3</i>		4	PDZ and LIM domain 3	NM_014476.5	DCM, HCM	
<i>SLC25A3</i>		12	Solute carrier family 25 (mitochondrial carrier, phosphate carrier) member 3	NM_145305.2		
<i>TNNT2</i>		1	Troponin T type 2 (cardiac)	NM_001276345.1	DCM, LVNC	
<i>TPM1</i>		15	Tropomyosin 1(alpha)	NM_000366.5	DCM, LVNC	
<i>TTN</i>		2	Titin	NM_001267550.2	DCM, limb-girdle muscular dystrophy	
<i>VCL</i>		10	Vinculin	NM_014000.2	DCM	
<b>LVNC</b>		<i>DTNA</i>	18	Dystrobrevin, alpha	NM_001390.4	DCM
<b>Cardiac ion channelopathies</b>						
<b>Afib</b>		<i>CAMK2G</i>	10	Calcium/calmodulin-dependent protein kinase II gamma	NM_172171.2	
	<i>GATA5</i>	20	GATA-binding protein 5	NM_080473.4	DCM	
	<i>GATA6</i>	18	GATA binding protein 6	NM_005257.5	ASD, tetralogy of fallot	
	<i>GJA5</i>	1	Gap junction protein alpha 5	NM_005266.6		
	<i>GJD4</i>	10	Gap junction protein delta 4	NM_153368.2		
	<i>KCNA5</i>	12	Potassium voltage-gated channel, shaker-related subfamily, member 5	NM_002234.3	ASD, LDS	
	<i>RANGRF</i>	17	RAN guanine nucleotide release factor	NM_016492.4	BrS	
	<i>SCN2B</i>	11	Na channel Auxiliar B subunit type 2	NM_004588.4	ASD, BrS	

<b>BrS</b>	<i>CACNA1C</i>	12	Calcium channel, voltage-dependent, L type, a 1C subunit	NM_199460.3		
	<i>CACNA2D1</i>	7	Calcium channel, voltage-dependent, a 2/d subunit 1	NM_000722.2	LDS, SQTS	
	<i>CACNB2</i>	10	Calcium channel, voltage-dependent, b 2 subunit	NM_000724.3		
	<i>GPD1L</i>	3	Glycerol-3-phosphate dehydrogenase 1-like	NM_015141.3	SIDS	
	<i>HCN4</i>	15	Hyperpolarization-activated cyclic nucleotide-gated potassium channel 4	NM_005477.2	SIDS, Afib	
	<i>HCN2</i>	19	Hyperpolarization activated cyclic nucleotide-gated potassium channel 2	NM_001194.3		
	<i>KCND3</i>	1	Potassium voltage-gated channel, Shal-related family, member 3	NM_004980.4	SIDS, Afib	
	<i>KCNE3</i>	11	Potassium voltage-gated channel, Isk-related family, member 3	NM_005472.4	paramyotonia congenita	
	<i>KCNJ8</i>	12	Potassium inwardly rectifying channel, subfamily J, member 8	NM_004982.3	ASD, LDS, SIDS	
	<i>SCN1B</i>	19	Na channel Auxiliari B subunit type 1	NM_199037.3	Afib	
	<i>SCN3B</i>	11	Na channel Auxiliari B subunit type 3	NM_018400.3	Afib, SIDS	
	<i>SCN4B</i>	11	Na channel Auxiliari B subunit type 4	NM_174934.3	Afib, SIDS	
	<i>SCN5A</i>	3	Cardiac voltage gated sodium channel	NM_001099404.1	ARVC, Afib, DCM, LQTS, SIDS	
	<i>SLMAP</i>	3	Sarcolemma associated protein	NM_001304420.2		
	<i>SCN10A</i>	3	Sodium channel type 10	NM_001293306.2	Afib,	
	<i>TRPM4</i>	19	Transient receptor potential cation channel, subfamily M, member 4	NM_017636.3	familial progressive cardiac conduction defect	
	<b>CPVT</b>	<i>CASQ2</i>	1	Calsequestrin 2 (cardiac muscle)	NM_001232.3	HCM
		<i>RYR2</i>	1	Ryanodine receptor	NM_001035.2	ARVC, LQTS
<i>TRDN</i>		6	Triadin	NM_006073.3	DCM, LQTS	
<b>IVF</b>	<i>ABCC8</i>	11	ATP-binding cassette, subfamily C (CFTR/MRP), member 8	NM_001287174.1	familial hyperinsulinemic hypoglycemia, hypoglycemia of infancy	
	<i>DPP6</i>	7	Dipeptidyl-peptidase 6	NM_130797.3		
	<i>MED23</i>	6	Mediator complex subunit 23	NM_004830.3	intellectual disability	
<b>LQTS</b>	<i>SEMA3A</i>	7	Sema domain, immunoglobulin domain (Ig), short basic domain secreted 3A	NM_006080.2	BrS, Kallmann syndrome	
	<i>AKAP9</i>	7	A kinase (PRKA) anchor protein (yotiao) 9	NM_001148.4	BrS, LDS	
	<i>ANK2</i>	4	Ankyrin 2	NM_001148.4		
	<i>CALM1</i>	14	Calmodulin 1	NM_006888.4	SIDS, CPVT	
	<i>CALM2</i>	2	Calmodulin 2	NM_001743.4	SIDS, CPVT	
	<i>CALM3</i>	19	Calmodulin 3	NM_005184.2	SIDS, CPVT	
	<i>DLG1</i>	3	Discs, large homolog 1	NM_001098424.1		
	<i>KCNE1</i>	21	Potassium voltage-gated channel, Isk-related family, member 1	NM_000219.5	Afib	
	<i>KCNE1L</i>	X	KCNE1-like	NM_012282.2	Afib	
	<i>KCNE2</i>	21	Potassium voltage-gated channel, Isk-related family, member 2	NM_172201.1		
	<i>KCNE5</i>	X	Potassium voltage-gated channel, subfamily E	NM_012282.2		
	<i>KCNH2</i>	7	Potassium Voltage gated channel, subfamily H, member 2	NM_000238.3		
	<i>KCNJ2</i>	17	Potassium inwardly rectifying channel, subfamily J, member 2	NM_000891.2	Afib, SQTS	
	<i>KCNJ5</i>	11	Potassium inwardly rectifying channel, subfamily J, member 5	NM_000890.3		
	<i>KCNQ1</i>	11	Potassium voltage gated channel, KQT type	NM_000218.2	Afib, SQTS	
	<i>NOS1AP</i>	1	Nitric oxide synthase 1 (neuronal) adaptor protein	NM_014697.2		
	<i>SNTA1</i>	20	Syntrophin alpha 1	NM_003098.2	BrS, LDS, SIDS	
	<b>Cardiovascular diseases</b>					
<b>AOVD</b>	<i>NOTCH1</i>	9	Notch 1	NM_017617.3		
<b>ASD</b>	<i>GATA4</i>	8	GATA-binding protein 4	NM_002052.3		
	<i>GATA6</i>	18	GATA-binding protein 6	NM_005257.5		
	<i>GJA1</i>	6	Gap dystroph protein alpha 1	NM_000165.4	SIDS	
	<i>Nkx2-5</i>	5	NK2 transcription factor-related 5	NM_004387.3	tetralogy of fallot	
	<i>TBX5</i>	12	T-box 5	NM_000192.3	Holt-oram sndrome	
<b>CVDX</b>	<i>FLNA</i>	X	Filamin A, alpha	NM_001110556.1		
<b>MVP</b>	<i>DCHS1</i>	11	Dachsous cadherin-related 1	NM_003737.3		
<b>Tricuspid atresia</b>	<i>HEY2</i>	6	Hes-related family BHLH transcription factor with YRPW motif 2	NM_012259.2		
<b>TAAD</b>	<i>ACTA2</i>	10	Actin, alpha 2, smooth muscle, aorta	NM_001141945.1	multisystemic smooth muscle dysfunction syndrome	
	<i>PRKG1</i>	10	Protein kinase, cGMP-dependent type 1	NM_006258.3	phosphoglycerate kinase deficiency	
<b>Connective tissue diseases</b>						
<b>WMS</b>	<i>ADAMTS10</i>	19	ADAM metallopeptidase with thrombospondin type 1 motif 10	NM_030957.3		
<b>EDS</b>	<i>COL3A1</i>	2	Collagen, type III, alpha 1	NM_000090.3		
	<i>COL5A1</i>	9	Collagen, type V, alpha 1	NM_000093.4		
	<i>COL5A2</i>	2	Collagen, type V, alpha 2	NM_000393.3		
<b>Cutis laxa</b>	<i>EFEMP2</i>	11	EGF containing fibulin-like extracellular matrix protein 2	NM_016938.4	AA, arteriopathy syndrome	
	<i>ELN</i>	7	Elastin	NM_001278939.1		
	<i>CTGF</i>	6	Connective tissue growth factor	NM_001901.2	HCM	
<b>MS</b>	<i>FBN1</i>	15	Fibrillin 1	NM_000138.4		
	<i>FBN2</i>	5	Fibrillin 2	NM_001999.3		
<b>LDS</b>	<i>TGFβ2</i>	1	Transforming growth factor beta 2	NM_001135599.2		
	<i>TGFβR1</i>	9	Transforming growth factor, beta receptor 1	NM_004612.2		
	<i>TGFβR2</i>	3	Transforming growth factor, beta receptor 2	NM_001024847.2		
	<i>SMAD3</i>	15	SMAD family member 3	NM_005902.3		
<b>Metabolic disease</b>						
<b>Fabry disease</b>	<i>GLA</i>	X	Galactosidase, alpha	NM_000169.2	HCM	
<b>GA</b>	<i>ETFA</i>	15	Electron-transfer-flavoprotein, alpha polypeptide	NM_001127716.1		
	<i>ETFB</i>	19	Electron-transfer-flavoprotein, beta polypeptide	NM_001985.2		
	<i>ETFDH</i>	4	Electron-transferring-flavoprotein dehydrogenase	NM_001281738.1		
<b>GKD</b>	<i>GK</i>	X	Glycerol kinase	NM_001128127.2		
	<i>GLB1</i>	3	Galactosidase, beta 1	NM_000404.2		
<b>GSD</b>	<i>AGL</i>	1	Amylo-alpha-1,6-glucosidase, 4-alpha-glucanotransferase	NM_000028.2		
	<i>G6PC</i>	17	Glucose-6-phosphatase, catalytic subunit	NM_000151.3		
	<i>GUSB</i>	7	Glucuronidase, beta	NM_000181.3		
	<i>SLC37A4</i>	11	Solute carrier family 37 (glucose-6-phosphate transporter)	NM_001164278.1		
<b>HHF</b>	<i>HADH</i>	4	Hydroxyacyl-CoA dehydrogenase	NM_001184705.2		
	<i>HADHA</i>	2	Hydroxyacyl-CoA dehydrogenase/3-ketoacyl-CoA thiolase/enoyl-CoA hydratase, alpha subunit	NM_000182.4		
	<i>HADHB</i>	2	Hydroxyacyl-CoA dehydrogenase/3-ketoacyl-CoA thiolase/enoyl-CoA hydratase, beta subunit	NM_001281512.1		
<b>HMGCS-deficiency</b>	<i>HMGCL</i>	1	3-hydroxymethyl-3-methylglutaryl-CoA lyase	NM_001166059.1		
	<i>HMGCS2</i>	1	3-hydroxy-3-methylglutaryl-CoA synthase 2 (mitochondrial)	NM_001166107.1		
<b>MCAD</b>	<i>ACADM</i>	1	Acyl-CoA dehydrogenase, C-4 to C-12 straight chain	NM_001127328.2		
	<i>ACAD9</i>	3	Acyl-CoA dehydrogenase family, member 9	NM_014049.4	muscular dystrophy	
	<i>ACADS</i>	12	Acyl-CoA dehydrogenase, C-2 to C-3 short chain	NM_001302554.1		
	<i>ACADVL</i>	17	Acyl-CoA dehydrogenase, very long chain	NM_001270447.1		
<b>mt-D</b>	<i>ATP5E</i>	20	ATP synthase, H+ transporting, mitochondrial F1 complex, epsilon subunit	NM_006886.3		
	<i>COA5</i>	2	Cytochrome c oxidase assembly factor 5	NM_001008215.2	fatal infantile cardioencephalomyopathy	
	<i>FOXRED1</i>	11	FAD-dependent oxidoreductase domain containing 1	NM_017547.3		
	<i>SCO2</i>	22	SCO2 cytochrome c oxidase assembly protein	NM_001169109.1		
	<i>SLC25A3</i>	12	Solute carrier family 25 (mitochondrial carrier, phosphate carrier) member 3	NM_145305.2	carnitine-acylcarnitine translocase deficiency	

<b>SPCD</b>	<i>CPT1A</i>	11	Carnitine palmitoyltransferase 1A (liver)	NM_001031847.2	
	<i>CPT2</i>	1	Carnitine palmitoyltransferase 2	NM_000098.2	
	<i>SLC22A5</i>	5	Solute carrier family 22 (organic cation/carnitine transporter), member 5	NM_001270888.1	
<b>Respiratory diseases</b>					
<b>CCHS</b>	<i>ASCL1</i>	12	Achaete-scute family bHLH transcription factor 1	NM_004316.3	
<b>PAH</b>	<i>ACVRL1</i>	12	Activin A receptor type II-like 1	NM_000020.2	
	<i>BMPRI1B</i>	4	Bone morphogenetic protein receptor, type IB	NM_001256793.1	
	<i>BMPRI2</i>	2	Bone morphogenetic protein receptor, type II	NM_001204.6	
	<i>CAVI</i>	7	Caveolin 1	NM_001753.4	
	<i>ENG</i>	9	Endoglin	NM_001114753.2	
	<i>SMAD9</i>	13	SMAd family member 9	NM_001127217.2	
<b>Muscular dystrophy</b>					
<b>SCD</b>	<i>ZNF365</i>	10	Zinc finger protein 365	NM_199451.2	
<b>CFC1</b>	<i>BRAF</i>	7	v-raf murine sarcoma viral oncogene homolog B	NM_004333.4	HCM
<b>NSLL</b>	<i>CBL</i>	11	Cbl proto-oncogene, E3 ubiquitin protein ligase	NM_005188.3	HCM
<b>DM1</b>	<i>DMPK</i>	19	Dystrophia myotonica-protein kinase	NM_001288764.1	DCM
<b>CMD</b>	<i>FKRP</i>	19	Fukutin related protein	NM_001039885.2	
<b>EDDM</b>	<i>SYNE1</i>	6	Spectrin repeat containing, nuclear envelope 1	NM_033071.3	
	<i>EMD</i>	X	Emerin (Emery-Dreifuss muscular dystrophy)	NM_000117.2	
	<i>SYNE2</i>	14	Spectrin repeat containing, nuclear envelope 2	NM_015180.4	
	<i>HFE</i>	6	Hemochromatosis	NM_000410.3	
	<i>ECE1</i>	1	Endothelin converting enzyme 1	NM_001113347.1	hirschsprung disease, cardiac defects, and autonomic dysfunction
<b>EMPF</b>	<i>DNM1L</i>	12	Dynammin 1-like	NM_001278464.1	

Abbreviations: AA: aortic aneurysm, Afib: atrial fibrillation, AOVD: aortic valve disease, ARVC: arrhythmogenic right ventricular cardiomyopathy, ASD: atrial septal defect, ATS: arterial tortuosity syndrome, BrS: Brugada syndrome, BTHS: Barth syndrome, CCHS: congenital central hypoventilation syndrome, CFC1: cardiofaciocutaneous syndrome 1, CMD: congenital muscular dystrophy, CPVT: catecholaminergic polymorphic ventricular tachycardia, CVDX: cardiac valvular dysplasia x-linked, DCM: dilated cardiomyopathy, DM1: myotonic dystrophy type 1, EDS: Ehlers-Danlos syndrome, EDMD: Emery-dreiss muscular dystrophy, EMPF: encephalopathy due to defective mitochondrial and peroxisomal fission, GA: glutaric aciduria, GKD: glycerol kinase deficiency, GSD: glycogen storage disease, HHF: familial hyperinsulinemic hypoglycaemia, HCM: hypertrophic cardiomyopathy, HMGCS: HMG-CoA synthase deficiency, IVF: idiopathic ventricular fibrillation, LDS: Loeys-Dietz syndrome, LQTS: long QT syndrome, LVNC: left ventricular noncompaction cardiomyopathy, MCAD: medium-chain-acyl-CoA-dehydrogenase deficiency, MS: Marfan syndrome, mt-D: mitochondrial defects, MVP: mitral valve prolapse, NSLL: Noonan syndrome-like disorder with or without juvenile myelomonocytic leukaemia, PAH: pulmonary hypertension, SCD: sudden cardiac death, SIDS: sudden infant death syndrome, SPCD: systemic primary carnitine deficiency, SQTS: short QT syndrome, SSS: sick sinus syndrome 2, TAAD: familial thoracic aortic aneurysm and aortic dissection, WMS: weill-marchesani syndrome.